## EXHIBIT H

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IN THE UNITED STATES DISTRICT COURT FOR THE SOUTHERN
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         DISTRICT OF WEST VIRGINIA-CHARLESTON DIVISION
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    _____
    IN RE: ETHICON, INC.
 5
    PELVIC REPAIR SYSTEM,
    PRODUCTS LIABILITY LITIGATION: MDL NO. 2327
 6
    ______
    THIS DOCUMENT RELATES TO ALL CASES
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9
          CONFIDENTIAL - SUBJECT TO PROTECTIVE ORDER
10
                        July 15, 2013
11
12
13
14
           Deposition of RAMY MAHMOUD, M.D., held
15
     at DRINKER BIDDLE AND REATH, LLP, 105 College
16
     Road East, Suite 300, Princeton, New Jersey,
17
     commencing at approximately 9:20 a.m., before
18
     Margaret M. Reihl, a Certified Realtime
19
     Reporter, Certified Court Reporter and Notary
20
     Public for the State of New Jersey.
21
22
23
                   GOLKOW TECHNOLOGIES, INC.
24
                877.370.3377 ph 917.591.5672 fax
                      deps@golkow.com
25
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- 1 have reported to your subordinates, I don't want to ask
- 2 you to name them all.
- A. No, it was a very small number. It
- 4 might have been perhaps three. I don't recall the
- 5 exact number.
- Q. Who were the people in Germany that
- 7 reported to you or your subordinates?
- A. I don't recall the names. I can tell
- 9 you that their function was preclinical research. At
- 10 one point there may also have been a medical affairs
- 11 physician in Germany and at other times not.
- 12 Q. The preclinical research that was
- 13 performed in Germany during your time with Ethicon, did
- it involve any of the transvaginal tapes or
- 15 transvaginal mesh?
- 16 A. I don't recall.
- 0. Go ahead and tell me what were your
- 18 responsibilities as chief medical officer of Ethicon
- 19 and vice president in charge of evidence-based
- 20 medicine?
- 21 A. So there were four departments that
- 22 comprised evidence-based medicine. One of them was
- 23 preclinical research. One of them was health economics
- 24 and reimbursement. One of them was clinical
- 25 development, and the last was medical affairs.

- Q. What was the third one? I'm sorry.
- 2 A. Clinical development.
- Q. And so all four of these departments
- 4 reported to you?
- 5 A. Yes, each of those departments had a
- 6 designated leader, and each of those leaders reported
- 7 to me.
- 8 Q. Who was the leader for preclinical?
- 9 A. Well, that changed over time.
- 10 Q. Okay.
- 11 A. When I first arrived it was a Dr. Mark
- 12 Storch, and then I later hired a new leader for
- 13 preclinical research. His name was Larry Johnson.
- Q. And who was the leader for health
- 15 economics and research?
- 16 A. Health economics and reimbursement.
- 17 Q. I'm sorry, reimbursement, excuse me.
- 18 A. For the majority of the time that I was
- 19 there, the leader was named Sheri Dodd.
- O. And who was the leader for clinical
- 21 development?
- 22 A. Jessica Shen.
- 0. Is that Cheng?
- 24 A. Shen, S-h-e-n.
- Q. And I believe you told me the medical

- 1 A. Yes.
- Q. Did you have an understanding during
- 3 your tenure or were you aware of this document hold?
- 4 A. Almost certainly. I didn't discard
- 5 documents pretty much at all. I operate under the
- 6 general assumption that nothing could be discarded
- 7 unless I knew specifically that it could.
- 8 O. And what was your process for that? I
- 9 mean, was there a policy that you were to discard or
- 10 destroy documents that were not subject to litigation
- 11 holds or some other sort of hold periodically?
- 12 A. There was a document retention policy
- for the company as a whole, to which I would have
- 14 adhered, but I cannot recite for you what that policy
- 15 was.
- Q. Was it your understanding that -- well,
- 17 back up and strike that.
- 18 Throughout your time at Ethicon, did you
- 19 retain all of your documents related to pelvic mesh
- 20 products, to the best of your knowledge?
- 21 A. I'm confident that I complied with the
- 22 retention policy, which included retaining all the
- 23 documents for which a document hold notice had been
- 24 issued.
- Q. And you understood or you operated under